PATENT COOPERATION TREATY PCT

INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY

(Chapter II of the Patent Cooperation Treaty)

(PCT Article 36 and Rule 70)

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Applicant's or agent's file reference 730815	FOR FURTHER ACTION	ee Form PCT/IPEA/416			
International application No.	International filing date (day/month/year)	Priority date (day/month/year)			
PCT/SG2004/000353	26 October 2004	27 October 2003			
International Patent Classification (IPC) or national classification and IPC					
Int. Cl. ⁷ C07C 275/50, 275/54, 311/58, 311/60, 335/26, C07D 209/14, 213/40, 233/61, 235/14, 285/06, 295/13, 295/215, 307/14, 307/85, 333/20, 333/38, 333/70, A61K 31/17, 31/341, 31/343, 31/44, 31/445, 31/381, 31/404, 31/433, 31/64, A61P 35/00					
Applicant S*BIO PTE LTD et al					
	ary examination report, established by this Inte ted to the applicant according to Article 36.	rnational Preliminary Examining			
2. This REPORT consists of a total of 5	sheets, including this cover sheet.				
3. This report is also accompanied by ANN	NEXES, comprising:				
a. X (sent to the applicant and to the	International Bureau) a total of 2 sheets, as	follows:			
x sheets of the description, c sheets containing rectificat Administrative Instruction	laims and/or drawings which have been amendions authorized by this Authority (see Rule 70s).	ded and are the basis for this report and/or .16 and Section 607 of the			
sheets which supersede earlier sheets, but which this Authority considers contain an amendment that goes beyond the disclosure in the international application as filed, as indicated in item 4 of Box No. I and the Supplemental Box.					
a sequence listing and/or table re	u only) a total of (indicate type and number of elated thereto, in computer readable form only be Section 802 of the Administrative Instruction	, as indicated in the Supplemental Box			
4. This report contains indications relating	to the following items:				
X Box No. I Basis of the report	t				
Box No. II Priority					
	of opinion with regard to novelty, inventive step and industrial applicability				
		seep and industrial applicationity			
X Box No. V Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement					
X Box No. VI Certain documents	X Box No. VI Certain documents cited				
Box No. VII Certain defects in	the international application				
Box No. VIII Certain observations on the international application					
Date of submission of the demand Date of completion of the report					
18 July 2005	23 September 2005	Date of completion of the report			
Name and mailing address of the IPEA/AU	Authorized Officer				
AUSTRALIAN PATENT OFFICE PO BOX 200, WODEN ACT 2606, AUSTRAL E-mail address: pct@ipaustralia.gov.au	S.R. IDRUS	AUD I			
Facsimile No. (02).6285 3929	Telephone No. (02) 62	83 2659			

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Box	No. I	Basis of the report					
1.		egard to the language, this report is based on the international application in the language in which it was filed, unless rise indicated under this item.					
	This report is based on translations from the original language into the following language, which is the language of a translation furnished for the purposes of:						
	international search (under Rules 12.3 and 23.1 (b))						
	publication of the international application (under Rule 12.4)						
	[international preliminary examination (under Rules 55.2 and/or 55.3)					
2.	With regard to the elements of the international application, this report is based on (replacement sheets which have been furnished to the receiving Office in response to an invitation under Article 14 are referred to in this report as "originally filed" and are not annexed to this report):						
	=	ne international application as originally filed/furnished					
	X th	ne description:					
		pages 1-3, 5-117 as originally filed/furnished					
		pages* 4 received by this Authority on 18 July 2005 with the letter of 15 July 2005 pages* received by this Authority on with the letter of					
	X th	e claims:					
		pages 118, 120-140 as originally filed/furnished					
		pages* as amended (together with any statement) under Article 19 pages* 119 received by this Authority on 18 July 2005 with the letter of 15 July 2005					
		pages* received by this Authority on with the letter of					
	. th	e drawings:					
		pages as originally filed/furnished					
•		pages* received by this Authority on with the letter of pages* received by this Authority on with the letter of					
	a	sequence listing and/or any related table(s) - see Supplemental Box Relating to Sequence Listing.					
3.	T	he amendments have resulted in the cancellation of:					
		the description, pages					
		the claims, Nos.					
		the drawings, sheets/figs					
		the sequence listing (specify):					
		any table(s) related to the sequence listing (specify):					
4 .	m:	nis report has been established as if (some of) the amendments annexed to this report and listed below had not been ade, since they have been considered to go beyond the disclosure as filed, as indicated in the Supplemental Box (Rule 0.2(c)).					
		the description, pages					
		the claims, Nos.					
		the drawings, sheets/figs					
		the sequence listing (specify):					
		any table(s) related to the sequence listing (specify):					
	16.						
	ıj uem	4 applies, some or all of those sheets may be marked "superseded."					

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Bo	x No.	III Non-	establishment of	opinion	with regard	to novelty,	inventive	step and i	ndustrial a	pplicabili	ty
1.	The questions whether the claimed invention appears to be novel, to involve an inventive step (to be non obvious), or to be industrially applicable have not been examined in respect of:										
,·		the entire int	ernational applica	ation						٠	
	X	claims Nos:	1-35 (in part)	•					·		
	beca	ause:						·			
		•	mational applicati								
		relate to the	following subject	matter w	hich does no	t require an	internation	al prelimin	ary examin	ation (spec	cify):
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	•										•
			on, claims or drav		=		· .	said claim	s Nos.		
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			·							-	
•		the claims, or	r said claims Nos.		•					•	
	-	are so inadeq	uately supported	by the des	scription that	no meaning	gful opinio	n could be	formed.		·
X no international search report has been established for said claim Nos. 1-35 (in part)											
	the nucleotide and/or amino acid sequence listing does not comply with the standard provided for in Annex C of the Administrative Instructions in that:								C of the		
	t	he written form	m	has	not been fur	nished		-			,
				doe	s not comply	with the st	andard				
	, t	he computer re	eadable form	has	not been fur	nished					
				doe	s not comply	with the st	andard	•		•	
		the tables rela	ated to the nucleo nical requirement	tide and/o s provided	r amino acid I for in Anne	sequence li ex C-bis of t	isting, if in he Admini	computer i	readable for tructions.	rm only, do	not comply
			ental Box for furt								

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Box No. V	Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability;
citation	s and explanations supporting such statement

1. Statement		·			
Novelty (N)	Claims 1-79	YES			
	Claims	NO			
Inventive step (IS)	Claims 1-79	YES			
	Claims	NO			
Industrial applicability (IA)	Claims 1-79	YES			
	Claims	NO			

2. Citations and explanations (Rule 70.7)

Where no international search report was established in respect of certain claims or parts thereof, the question whether the claimed invention is novel, involve inventive step and industrially applicable have not been examined.

For the search that was conducted, the International Search Report identified the following citations:

- D1) STN File CA Abstract Accession No. 133:53160
- D2) STN File CA Abstract Accession No. 132:342787
- D3) Derwent Abstract Accession No. 2004-383257/36

D1 and **D2** are the closest prior-art and disclosed compounds which are excluded by Claims 1-15, and 18. As such there is no overlap between the subject matter disclosed in the prior art documents and the subject matter of the present claims. Moreover, there is no teaching or suggestion in **D1** and **D2** that the compounds of the prior art would be useful for the inhibition of histone deacetylase and, as such, could be used for the treatment of cancer.

Accordingly, the claimed subject matter are novel and involve inventive step in the light of D1 and D2.

D3 is an intermediate document and is mentioned in Box VI.

The compounds of the present invention are inhibitors of histone deacetylase (HDACs) and it is understood that inhibition of HDACs will cause apoptosis of cancer cells. Thus the claimed subject matter has industrial applicability.

With regard to the document(s) listed in Box VI under "certain documents cited", these are documents published prior to the international filing date but later than the priority date claimed but which would otherwise be considered to be of particular relevance.

Under the PCT, novelty is considered only in respect of documents published before the priority date. The relevance of a document published after the priority date is dependent upon national law. Such documents are excluded from consideration in preliminary examination, under the PCT Guidelines but have been included here for information.

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Box No. VI Certain documents	cited		
1. Certain published documents (Rule	70.10)		
Application No. Patent No.	Publication date (day/month/year)	Filing date (day/month/year)	Priority date (valid claim)(day/month/year)
JP 2004143053 A ("X")	20 May 2004	22 October 2002	22 October 2002
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	:	•	
			•
			·
This application disclosed compount C=O which would be equivalent to I Q in present claims and R ² is equivalent 1-16, and 18.	M; L is G ² NH so that	it's C(=O)NH wherein the	e C(=O) would be equivalent to the
		•	
	·		
2. Non-written disclosures (Rule 70.9)			
Kind of non-written disclosure		vritten disclosure onth/year)	Date of written disclosure referring to non-written disclosure (day/month/year)
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heterocycloalkylalkyl, arylalkyl, heteroarylalkyl and acyl; each of which may be optionally substituted;

or a pharmaceutically acceptable salt or prodrug thereof, wherein when R is methyl or isopropyl methyl then R₂ is not benzyl.

In one preferred embodiment the present invention provides compounds having the Formula (2)

$$G - Q - N - N - B - A - Z - L - O$$

$$R^{1}$$

$$N - B$$

$$R^{2}$$

$$N - OH$$

Formula (2)

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wherein

R¹ is selected from the group consisting of H, C₁ -C₆ alkyl and acyl;

L is a single bond or is a C_1 - C_5 hydrocarbon chain which may contain 0 to 2 multiple bonds independently selected from double bonds and triple bonds and wherein, the chain may optionally be interrupted by at least one of -O-, -S-, -S(O)- and -S(O)₂- and the chain may optionally be substituted with one or more substituents independently selected from the group consisting of C_1 - C_4 alkyl;

Z is selected from the group consisting of a single bond, $N(R^1)$, O, S, S(O) and $S(O)_2$;

A is selected from the group consisting of a single bond, optionally substituted arylene, optionally substituted heteroarylene, optionally substituted cycloalkylene and optionally substituted heterocycloalkylene;

B is selected from the group consisting of a single bond, optionally substituted aminoacyl, optionally substituted arylene, optionally substituted heteroarylene, optionally substituted arylalkylene, optionally substituted heteroarylene, optionally substituted alkylene, optionally substituted C_1 - C_3 alkylene, optionally substituted heteroalkylene, optionally substituted cycloalkylene, optionally substituted heterocycloalkylene and optionally substituted - $(CH_2)_m$ -C(O)- $N(R^4)$ - $C(CH_2)_n$ -, wherein n is an integer from 0 to 6, m is an integer from 0 to 6;

heteroaryloxy, arylalkyloxy, amino, alkylamino, aminoalkyl, acylamino, arylamino, sulfonylamino, sulfinylamino, phenoxy, benzyloxy, COOR⁴, CONHR⁴, NHCOR⁴, NHCOOR⁴, NHCONHR⁴, C(=NOH)R⁴, alkoxycarbonyl, alkylaminocarbonyl, sulfonyl, alkylsulfonyl, arylsulfonyl, arylsulfinyl, aminosulfonyl, aminosulfinyl, SR⁴ and acyl; each of which may optionally be substituted;

Q is selected from the group consisting of $-S(O)_2$ -, -C(=O)- and -C(=S)-;

G is selected from the group consisting of optionally substituted alkyl, optionally substituted cycloalkyl, optionally substituted aryl, optionally substituted heteroaryl, optionally substituted heterocycloalkyl, optionally substituted arylalkyl, and optionally substituted heteroarylalkyl;

each R⁴ is independently selected from the group consisting of H, alkyl, alkenyl, alkynyl, haloalkyl, heteroalkyl, cycloalkyl, heterocycloalkyl, aryl, heteroaryl, cycloalkylalkyl, heterocycloalkylalkyl, arylalkyl, heteroarylalkyl and acyl, each of which may be optionally substituted;

or a pharmaceutically acceptable salt or prodrug thereof, wherein when R is methyl or isopropyl methyl then R₂ is not benzyl.

2. A compound according to claim 1 having the Formula (2)

25 wherein

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R¹ is selected from the group consisting of H, C₁ -C₆ alkyl and acyl;

L is a single bond or is a C_1 - C_5 hydrocarbon chain which may contain 0 to 2 multiple bonds independently selected from double bonds and triple bonds and wherein, the chain may optionally be interrupted by at least one of -O-, -S-, -S(O)- and -S(O)₂- and the chain may optionally be substituted with one or more substituents independently selected from the group consisting of C_1 - C_4 alkyl;